

## O C C A S I O N A L I D E A S

# Twelve Common Questions About Human Rabies and Its Prevention

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Rabies, described as early as the 23rd century b.c., may be one of the oldest recorded infectious diseases of mankind [1]. It is a fatal infection caused by a highly neurotropic, bullet-shaped, single-stranded RNA virus [2]. Rabies causes more than 40,000–60,000 deaths worldwide each year, primarily in countries in which canine rabies is endemic and the delivery of health care is difficult [3,4]. In the United States, human rabies is rare; just 27 deaths have occurred since 1990 [5,6]. Nevertheless, the scarcity of cases of human rabies does not minimize its importance to public health nor the fear it instills in individual patients who are potentially exposed.

The epidemiology of human rabies is ultimately linked to cycles of rabies virus transmission in animals. The low rate of human infection is due to animal control, animal vaccination programs, and the effective application of human pre-exposure and postexposure prophylaxis. In combination, these efforts cost more than \$300 million annually [7], and each year between 16,000 and 39,000 people receive postexposure prophylaxis in the United States [8]. This article addresses some common questions about rabies and its prevention.

## 1

*Before departing for rural Latin America to study, a graduate student questions whether she should receive primary rabies vaccination.*

**Who should receive the primary or preexposure vaccination?**

Preexposure prophylaxis, given by the intradermal (ID) or intramuscular (IM) route, is indicated for certain high-risk groups (Table 1) [9]. The decision to give preexposure prophylaxis to travelers is complicated by several factors: a very low incidence of rabies reported in travelers, the relatively high cost of preexposure rabies prophylaxis (in excess of \$300), and the need to begin the vaccination series at least 3 weeks before departure [10]. Nevertheless, preexposure prophylaxis should be considered for international travelers who are likely to come in contact with animals in areas where canine rabies is present and where immediate access to appropriate medical care, including safe and effective biologic agents, may be difficult. One source of information on rabies in various geographic regions is the Centers for Disease Control and Prevention Travelers' Health website [11].

Preexposure prophylaxis simplifies postexposure prophylaxis by eliminating the need for rabies immunoglobulin (RIG) and may provide a measure of protection in the event that a true exposure is not recognized. It does not eliminate the need for appropriate wound treatment and additional vaccinations in the case of a known exposure. Routine serologic testing to confirm seroconversion is not necessary except in the case of immunocompromised individuals. Chloroquine has been noted to reduce the patient's response to the vaccine. Therefore, if the patient receives ID administration of preexposure prophylaxis, it must be completed before the initiation of antimalarial treatment. If the patient is already taking antimalaria medications, he or she should receive IM administration of the vaccine [9].

## 2

*A missionary traveling to Southeast Asia to perform health care work wonders whether the rabies vaccine is available should he need it.*

**What types of rabies biologic agents are available abroad?**

Cell culture vaccines are clearly the standard; they are effective and well tolerated. In the United States, human diploid cell vaccine (HDCV), rabies vaccine adsorbed (RVA) derived from fetal rhesus lung diploid cell, and purified chick embryo-cell vaccine (PCEC) are available. Other cell culture vaccines used outside the United States include purified Vero-cell rabies vaccine and purified duck-embryo vaccine. In addition, many countries still use vaccines produced from animal nerve tissue that have a high rate of adverse reactions

**TABLE 1. Rabies preexposure prophylaxis guide, United States, 1999**

Risk category	Nature of risk	Typical populations	Preexposure recommendations
	Virus present continuously, often in high concentrations; specific exposures likely to go unrecognized; bite, nonbite, or aerosol exposure	Rabies research laboratory workers <sup>a</sup> ; rabies biologic agents production workers	Primary course; serologic testing every 6 mo; booster vaccination if antibody titer is below acceptable level <sup>b</sup>
Frequent	Exposure usually episodic, with source recognized, but exposure may also be unrecognized; bite, nonbite, or aerosol exposure	Rabies diagnostic laboratory workers, <sup>a</sup> spelunkers, veterinarians and staff, and animal-control and wildlife workers in rabies enzootic areas	Primary course; serologic testing every 2 y; booster vaccination if antibody titer is below acceptable level <sup>b</sup>
Infrequent (greater than the population at large)	Exposure nearly always episodic with source recognized; bite or nonbite exposure	Veterinarians and animal-control and wildlife workers in areas of low rabies rates; veterinary students; travelers visiting areas where rabies is enzootic and immediate access to appropriate medical care including biologic agents is limited	Primary course; no serologic testing or booster vaccination
Rare (population at large)	Exposures always episodic with source recognized; bite or nonbite exposure	U.S. population at large, including persons in rabies epizootic areas	No vaccination necessary

<sup>a</sup> Judgment of relative risk and extra monitoring of vaccination status of laboratory workers is the responsibility of the laboratory supervisor (9).

<sup>b</sup> Minimum acceptable antibody level is complete virus neutralization at a 1:5 serum dilution ratio by rapid fluorescent focus inhibition test. A booster dose should be administered if the titer falls below this level.

(neuromuscular reactions in 1 per 200–2000 persons vaccinated). Purified equine RIG has been used effectively in countries where human RIG is not available. Travelers should be aware that in some areas, unpurified antirabies serum of equine-origin immunoglobulin may be offered and is associated with a high rate of serious adverse reactions, including anaphylaxis [3,9].

### 3

*A pet owner asks if she should obtain rabies vaccination because she believes saliva was transferred to her hands while playing with her dog that had killed a rabid raccoon earlier that day.*

What constitutes a rabies exposure?

Rabies is transmitted when the virus is introduced into bite wounds, through breaks in the skin, or onto mucous membranes. Three questions to ask are the following: (1) Was the person bitten?; (2) Did

saliva or central nervous system material from a potentially rabid animal contaminate an open wound or mucous membrane?; and (3) Was the animal in question a bat? (see question 4). If all can be answered "No," then no exposure occurred and postexposure prophylaxis is not required. Petting a rabid animal and coming into contact with blood, urine, or feces of a rabid animal do not constitute exposures and are not indications for prophylaxis [9]. If the answer to at least one of these questions is "Yes," then exposure to rabies is a possibility, and the likelihood that the animal has rabies must be considered (Table 2) [9].

Questions often arise regarding contact with saliva. In general, if the material containing the virus is dry, it can be considered noninfectious; the rabies virus is inactivated by desiccation and ultraviolet radiation. Rabies cannot be transmitted from an animal that does not have active infection. Infectious material (saliva or neural tissue) would have to get in the mouth or on the claw

of that animal (the dog in the above scenario) and then promptly introduced through the skin or onto mucous membranes. This would be only remotely feasible in an immediate sequential exposure, and such a case has never been described. Local and state health departments can assist in deciding the likelihood that an animal has rabies and whether postexposure prophylaxis is indicated. Animals exposed to rabid animals need to be evaluated by a veterinarian and reported to the local health department immediately [12].

### 4

*Waking their 16-month-old son in the morning, parents find a bat in the corner of the room and they question whether the child needs to be seen by a doctor.*

Why is there a concern about bats?

Excluding dog bites that occurred outside of the country, 22 (71%) of

the 31 human cases of rabies in the United States since 1980 have been associated with bat rabies virus variants. Although the histories sometimes conflict, of the 22 patients with rabies, only two reported a bat bite, 10–12 had apparent contact with bats (many with which saliva, teeth, or other contact likely occurred), and 8–10 reported no exposure to bats or other sources of infection [6,9]. In these latter cases, an unreported or undetected bat bite remains the most plausible hypothesis. Therefore, postexposure prophylaxis should be considered when direct contact between a person and a bat might have occurred, unless the person can be certain that a bite, scratch, or mucous membrane exposure did not occur; for example, consider the potential for direct contact in situations in which a sleeping person wakes to find a bat in the room or an adult witnesses a bat in the room with a previously unattended child, mentally disabled person, or intoxicated person [9]. Examination for a bite wound alone is inadequate [13]. If, as in the example above, the bat in question is available for testing, and a clear exposure has not occurred, initiation of postexposure prophylaxis

may be delayed 24–48 hours for testing of the bat. If a bite or other clear exposure occurs, postexposure prophylaxis should begin immediately and may be discontinued if laboratory testing shows that the bat is not rabid [14].

## 5

*Camping scouts notice several bats fluttering in the trees above them. Their adult leader wonders whether they should receive treatment.*

Can rabies be transmitted by aerosol transmission?

Although nonbite routes of infection are possible, they are exceedingly rare and not applicable to usual public exposures. There have been two reports of rabies transmission to laboratory workers. Both workers were exposed to concentrated aerosols of the rabies virus [15]. In the 1950s, two cases of purported aerosol transmission to humans were associated with Frio Cave, Texas, which was inhabited by millions of bats. Although the patients did not recall a bite before they died, complete exclusion of a bite is not possible [16]. The conditions under which aerosol transmission might occur

are rare and unique, and it would be extremely unusual for them to be applicable to public exposures. Merely seeing a bat or being in the vicinity of bats does not constitute an exposure.

## 6

*A nurse calls asking whether she should administer postexposure prophylaxis intradermally to a patient bitten by a rat.*

How is secondary or postexposure prophylaxis administered?

Rabies prophylaxis in the setting of exposures to small mammals, including rodents, lagomorphs (rabbits and hares), and insectivores (e.g., shrews), is almost never required (Table 2). These animals are not reservoirs, and there have been no documented cases of rabies transmission to humans by these animals [9]. If there is a question, a call to the local or state health department may be helpful.

When postexposure prophylaxis is required, all doses of the vaccine are given in the deltoid area by IM administration (anterolateral thigh is acceptable for small children). The gluteal area should not be

TABLE 2. Rabies postexposure prophylaxis guide, United States, 1999

Animal type	Evaluation and disposition of animal	Postexposure prophylaxis recommendations
	Healthy and available for 10 d of observation	Should not begin prophylaxis unless animal develops clinical signs of rabies <sup>a</sup>
	Rabid or suspected rabid	Immediately vaccinate
	Unknown (e.g., escaped)	Consult public health officials
Skunks, raccoons, foxes, and most other carnivores; bats	Regard as rabid unless animal is proven negative by laboratory tests <sup>b</sup>	Consider immediate vaccination
Livestock, small rodents, lagomorphs (rabbits and hares), large rodents (woodchucks and beavers), and other mammals	Consider individually	Consult public health officials; bites of squirrels, hamsters, guinea pigs, gerbils, chipmunks, rats, mice, other small rodents, rabbits, and hares almost never require antirabies postexposure prophylaxis

<sup>a</sup> During the 10-day observation period, begin postexposure prophylaxis at first sign of rabies in a dog, cat, or ferret that has bitten someone. If the animal exhibits clinical signs of rabies, it should be euthanized immediately and tested.

<sup>b</sup> The animal should be euthanized and tested as soon as possible. Holding for observation is not recommended. Discontinue vaccine if immunofluorescence test results of the animal are negative.

used. For those who have had prior vaccination (with HDCV, RVA, or PCEC, or who have a documented history of antibody response to other vaccines), only two doses are required and RIG should not be used. For those who did not receive a vaccine previously, RIG should be thoroughly infiltrated around the wound area. Any remaining RIG should be injected by IM administration at a site distant from where the vaccine was administered. If RIG was not given when vaccination was begun, it can be administered through the seventh day after vaccination was started. Beyond the seventh day, it is not indicated (Table 3). Although no controlled trials have been performed, extensive experience from many parts of the world indicates that postexposure prophylaxis, consisting of local wound treatment, passive immunization (RIG), and vaccination, is effective if given in an appropriate and timely manner [9].

## 7

*A student is bitten by a neighbor's dog on the way to school and presents to the emergency department for wound treatment. The emergency department providers wonder whether a rabies vaccination is needed at this time.*

When can rabies prophylaxis be delayed while animals are held for quarantine and observation?

A healthy dog, cat, or ferret that bites a person should be confined and observed for 10 days. Any illness in the animal should be reported immediately to the local health department and evaluated by a veterinarian. If the animal remains healthy, patients do not need to begin rabies prophylaxis. If the biting dog, cat, or ferret is a stray animal, it should either be observed for 10 days or be euthanized immediately and submitted for rabies

examination. The management of animals other than dogs, cats, and ferrets depends on the species, the circumstances of the bite, the epidemiology of rabies in the area, the biting animal's history and current health status, and the potential for exposure to rabies. Public health and animal control officials can assist in these actions and decisions [9,12].

## 8

*A physician calls the state health department inquiring whether she should request rabies diagnostic tests on a comatose patient.*

When should a diagnosis of rabies be considered, and what specimens should be sent?

Rabies should be considered in the differential diagnosis of any patient who presents with acute progressing encephalopathy of unknown

TABLE 3. Rabies postexposure prophylaxis schedule, United States, 1999

Vaccination status	Treatment	Regimen <sup>a</sup>
Not previously vaccinated	RIG	All postexposure treatment should begin with immediate thorough cleansing of all wounds with soap and water; if available, a virucidal agent such as povidone-iodine solution should be used to irrigate the wounds
	Vaccine	Administer 20 IU/kg body weight; if anatomically feasible, the full dose should be infiltrated around the wound(s), and any remaining volume should be administered IM at an anatomical site distant from vaccine administration; also, RIG should not be administered in the same syringe as vaccine; because RIG may partially suppress active production of antibody, no more than the recommended dose should be given
Previously vaccinated <sup>d</sup>	Wound cleansing	HDCV, RVA, or PCEC 1.0 mL IM (deltoid area <sup>b</sup> ), one each on days 0 <sup>c</sup> , 3, 7, 14, and 28.
	RIG	All postexposure treatment should begin with immediate thorough cleansing of all wounds with soap and water; if available, a virucidal agent such as povidone-iodine solution should be used to irrigate the wounds
	Vaccine	RIG should not be administered
		HDCV, RVA, or PCEC 1.0 mL IM (deltoid area <sup>b</sup> ), one each on days 0 <sup>c</sup> and 3.

Note. Abbreviations used: HDCV, human diploid cell vaccine; PCEC, purified chick embryo-cell vaccine; RIG, rabies immunoglobulin; RVA, rabies vaccine adsorbed; IM, intramuscular.

<sup>a</sup> These regimens are applicable for all age groups, including children.

<sup>b</sup> The deltoid area is the only acceptable site of vaccination for adults and older children. For younger children, the outer aspect of the thigh may be used. Vaccine should never be administered in the gluteal area.

<sup>c</sup> Day 0 is the day the first dose of vaccine is administered.

<sup>d</sup> Any person with a history of preexposure vaccination with HDCV, RVA, or PCEC; prior postexposure prophylaxis with HDCV, RVA, or PCEC; or previous vaccination with any other type of rabies vaccine and a documented history of antibody response to the prior vaccination.

cause. The lack of an exposure history should not deter pursuing the diagnosis, since most patients in the United States have no definitive exposure history [5]. Once symptoms of rabies begin, the natural history is rapid clinical deterioration and death. Patients with encephalopathy who are clinically improving generally do not need rabies testing.

If rabies is suspected, samples that should be sent for study include nuchal skin biopsy, saliva, serum, and cerebrospinal fluid. The postmortem diagnosis of rabies is determined by an examination of brain tissue. Because of the rarity of the disease and lack of effective treatment, an antemortem brain biopsy is not indicated. However, biopsies performed for other diseases (e.g., herpesvirus encephalitis), if negative, can be tested for evidence of rabies virus infection [2].

## 9

*An emergency department doctor sees a patient with an infected raccoon bite 6 weeks after the bite occurred and questions whether it is still useful to administer rabies prophylaxis.*

How long after an exposure would prophylaxis still be considered?

Treatment with RIG and the vaccine should be given immediately if the exposure was of high risk. The usual incubation period for rabies in humans is 3–8 weeks [4,17]. However, incubation periods of 6 years or longer have been documented [18]. Thus, if a true exposure has occurred, prophylaxis (including RIG) should still be given, regardless of the length of time between exposure and clinical presentation for evaluation, although no studies on the efficacy of such treatment are available.

## 10

*A mother whose daughter has been scratched by a stray cat wonders what the risk of the RIG and vaccine is for her child.*

What are the adverse reactions of the RIG and vaccine?

With HDCV, local reactions (pain, erythema, swelling, itching) have been commonly reported (30%–74%). Systemic reactions (headache, dizziness, abdominal pain, nausea, muscle aches) have been reported in 5%–40% of recipients. Three cases of neurologic illness resembling Guillain-Barré syndrome that resolved without sequelae within 12 weeks have been reported. Rare reports of other nervous system disorders have been temporally associated with the HDCV vaccine, but a causal relationship has not been established. There is no evidence that any viruses have ever been transmitted by commercially available RIG in the United States [9].

Of note, rabies is almost always caused by a bite; nonbite exposures rarely cause rabies. Local or state public health officials should be contacted about the prevalence of rabies in any given area and the need to administer prophylaxis for scratches or other low-risk encounters.

## 11

*A patient with HIV infection wonders whether his response to postexposure vaccination will be appropriate.*

Do immunosuppressed patients require special consideration?

Patients who are immunosuppressed should postpone preexposure vaccinations and avoid situations for which rabies preexposure

prophylaxis is indicated. If this is not possible, they should be vaccinated by IM injection and have their antibody titer checked 2–4 weeks after the series. When post-exposure prophylaxis is needed, it is also important that a serum sample be tested for rabies antibody to ensure that an acceptable response has developed. The minimum acceptable antibody level is complete virus neutralization at a 1:5 serum dilution ratio by the rapid fluorescent focus inhibition test [9]. No rabies postexposure failures have been attributed to HIV infection.

## 12

*A camper has had the first two vaccine doses of the postexposure series when the state laboratory reports that the skunk that bit him tested negative. He wonders whether he should continue the series.*

What is the value of a negative direct fluorescent antibody test?

The direct fluorescent antibody (DFA) test for the detection of the rabies virus antigen in brain tissue is used as the primary diagnostic test in every public health laboratory in the United States and has a sensitivity approaching 100% [2]. Rabies diagnosis and prophylaxis of potential human exposures are based on the observation that the rabies virus reaches the salivary glands and is excreted in saliva only after replication in the central nervous system. The absence of the rabies virus antigen in the brain of an animal as determined by DFA examination (i.e., a negative diagnostic test result) essentially precludes the presence of the virus in saliva, the risk of rabies transmission, and the need for postexposure prophylaxis. Since the initiation of current testing procedures

in 1958, there is no evidence that a false-negative laboratory test has ever led to rabies in a person who was subsequently left untreated [19].

## Summary

Current public health practices have made cases of human rabies rare in the United States. Preexposure prophylaxis is available for select high-risk groups. Potential exposures to rabies will continue to occur, and patients will often present to their physicians for guidance. A careful history of exposure and, if needed, consultation with state health departments can guide the use of post-exposure prophylaxis. Appropriate wound treatment and administration of rabies vaccines and RIG prevent rabies infection in exposed individuals. Further information, including the recommendations of the Advisory Committee on Immunization Practices, can be found at the following CDC website: [www.cdc.gov/ncidod/dvrd/rabies](http://www.cdc.gov/ncidod/dvrd/rabies).

## Disclaimer

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